INDEX

VOLUME 21 1966

A

Administering that Ounce of Prevention: New Drugs and Nuclear Reactors. Sept., p. 458; Oct., p. 488.

Advantages of, and Need for the Establishment of Uniform Guiding Principles and Model Standards for Food (The). Dec., p. 680.

Advertising

Current Problems in Food and Drug Advertising. Mar., p. 161.

Allergic Responses

New Jersey minority rule. May, p. 293.

Appraisal of Progress in Drug Marketing. Jan., p. 33.

Artificial Sweetners—Their Impact on the Food Laws. Feb., p. 111.

Atomic Energy Commission

Nuclear reactors. Oct., p. 488.

B

Ballard, Robert W.—Progress in Research—A Question. Jan., p. 28.

Buatti, Anthony T.—An Appraisal of Progress in Drug Marketing. Jan., p. 33.

Burditt, George M.

Packaging Industry Looks at the Model Law and Regulation (The). Aug., p. 396.

Uniformity—A Legislator's Viewpoint. Mar., p. 145.

C

Canada

Food safety. May, p. 283.

Cavers, David F.—Administering that Ounce of Prevention: New Drugs and Nuclear Reactors. Sept., p. 458; Oct., p. 488.

Chapman, R. A.—Food Safety in Canada. May, p. 283.

Clinical Evaluation of Drug Safety. June, p. 348.

Codex Alimentarius Commission

The Future of the Codex Alimentarius Commission. Apr., p. 201.

Third Session of the Joint FAO/WHO Codex Alimentarius Commission. Mar., p. 168.

Color Additives

Distinction between artificial and natural color. June, p. 322.

Common Market and Harmonization of the Food Laws (The). Sept., p. 440.

Compliance with Drug Abuse Control Amendments of 1965. July, p. 360.

Condon, William J.—Product Liability—1965. Mar., p. 154.

Congressional Investigations: Some Observations. Jan., p. 40.

Consumer Education

FDA's responsibility. July, p. 368.

FDA's role. Feb., p. 78.

Role of industry and government. June, p. 322.

Consumer Interest

President's message. Mar., p. 179.

Rights of the consumer. Feb., p. 92.

What the Consumer Expects, and Receives, from the Administration of the Federal Food, Drug and Cosmetic Act. July, p. 377.

Consumer Protection

Food industry. Jan., p. 6.

Consumer's Interest (The). Feb., p. 92.

Cook, C. W.—What the Food Industry Needs from the FDA. July, p. 373.

Coulston, Frederick—Toxicologic Aspects of Drug Safety. June, p. 336.

Crowley, Cornelius D.—Current Good Manufacturing Practice. Mar., p. 137.

Current Good Manufacturing Practice. Mar., p. 137.

Current Problems in Food and Drug Advertising. Mar., p. 161.

D

Davies, J. H. V.—The Future of the Codex Alimentarius Commission. Apr., p. 201.

Davis, A. D.—Food and Drug Administration Plans and Programs. Jan., p. 57.

Deceptive Packaging of Foods
FDA's power. Apr., p. 205.
FTC's power. May, p. 248.

Depew, Franklin M.
Introductory Statement (Twenty-first Annual Meeting of the Section on Food, Drug and Cosmetic Law of the New York State Bar Association). Feb., p. 109.
Report of the Third Session of the Joint FAO/WHO Codex Alimentarius Commission. Mar., p. 168.

Dougherty, James R.—Current Problems in Food and Drug Advertising. Mar., p. 161.

Drug Abuse Control Amendments of 1965
Compliance with Drug Abuse Control Amendments of 1965. July, p. 360.

Drug Amendments of 1962
Drug investigation. Apr., p. 184.

Drug Investigation
A lawyer's view. Apr., p. 184.

Drug Safety
Clinical evaluation. June, p. 348.
FDA. Feb., p. 68.
Toxicologic aspects. June, p. 336.

Drug Safety and the FDA. Feb., p. 68.

E

Edible Oils and Fats
Latin-American Food Code—Chap. VII. June, p. 312.

F

FDA Rules of Practice and Procedure
A Look at FDA's New Rules of Practice — and Problems Still Unsolved. Sept., p. 448.

FDA Statement and an Industry Response. July, p. 368.

Factory Inspection Under the Federal Food, Drug and Cosmetic Act (Section 704). Dec., p. 673.

Fair Packaging and Labeling Act Analysis. Dec., p. 632.

Federal Food, Drug & Cosmetic Act
Current Good Manufacturing Practice. Mar., p. 137.
Factory inspection. Dec., p. 673.

Federal Regulation
Proprietary drug industry. Aug., p. 431.

Federal Trade Commission
Deceptive packaging of foods. May, p. 248.

Food Additives
Developments since the Food Additives Amendment of 1958. Nov., p. 616.
Packaging Industries and the Food Additives Amendment of 1958. Dec., p. 647.
Paper cups and plates. June, p. 322.
Synthetic. June, p. 322.

Food Additives Amendment of 1958
Developments since the Amendment. Nov., p. 616.
Packaging industries. Dec., p. 647.

Food and Drug Administration
Deceptive Packaging of Foods. Apr., p. 205.
Economic adulteration of foods. Oct., p. 533; Nov., p. 552.

Factory inspection. May, p. 275.
Industry information programs. Jan., p. 52.
New drugs. Sept., p. 458.
Planning operations. Jan., p. 57.
Recent legal developments. Oct., p. 517.
Role in consumer education. Feb., p. 78.
Role in drug safety. Feb., p. 68.
Rules of Practice and Procedure. Sept., p. 448.
Status of FDA's litigation. June, p. 322.
What the Consumer Expects, and Receives, from the Administration of the Federal Food, Drug and Cosmetic Act. July, p. 377.
What the Food Industry Needs from the FDA. July, p. 373.

Food and Drug Administration and the Economic Adulteration of Foods (The). Oct., p. 533; Nov., p. 552.

Food and Drug Administration, Federal Trade Commission and the Deceptive Packaging of Foods. Apr., p. 205; May, p. 248.

Food and Drug Administration Industry Information Programs. Jan., p. 52.

Food and Drug Administration Plans and Programs. Jan., p. 57.

Food Industry and Consumer Protection (The). Jan., p. 6.

Food Package Labeling — Legal Requirements and Commercial Practices. Aug., p. 392.

Food Research and the Food Laws. Apr., p. 191.

Food Safety
Canada. May, p. 283.
Food Safety in Canada. May, p. 283.

Food Standards
Food Research and the Food Laws. Apr., p. 191.
Review. June, p. 322.

Foods for Regimens
Latin-American Food Code—Chap. XVIII. Aug., p. 404.

Forte, Wesley E.
Food and Drug Administration and the Economic Adulteration of Foods (The). Oct., p. 533; Nov., p. 552.
Food and Drug Administration, Federal Trade Commission and the Deceptive Packaging of Foods. Apr., p. 205; May, p. 248.

Freedman, Warren
A Hatband and a Tube of Lipstick: The New Jersey Minority Rule on Allergic Responses. May, p. 293.
Prescription or Ethical Drugs: Fallacies as to Warranties, Failure to Warn, and Strict Liability in Tort. Nov., p. 599.

Future of the Codex Alimentarius Commission (The). Apr., p. 201.

G

Gamble, Theodore R.—The Food Industry and Consumer Protection. Jan., p. 6.

"Generic Every Time" Case: Prescription Drug Industry *in Extremis* (The). Apr., p. 226.

Generic Name
The "Generic Every Time" Case: Prescription Drug Industry *in Extremis*. Apr., p. 226.

Goddard, James L.—FDA Statement and an Industry Response. July, p. 368.

Gould, Wilbur A.—Food Research and the Food Laws. Apr., p. 191.

Graham, Andrew J.—The Law Governing FDA Factory Inspection. May, p. 275.

H

Hammel, Robert W.—Physician Owned Pharmaceutical Companies—A Wrong Without a Remedy. Aug., p. 412.

Harmonization
Common Market and Harmonization of the Food Laws. Sept., p. 440.
National food laws. Dec., p. 680.

Hatband and a Tube of Lipstick: The New Jersey Minority Rule on Allergic Responses (A). May, p. 293.

Heckman, Jerome H.—The Packaging Industries and the Food Additives Amendment of 1958—It's Time for a Change in the Law. Dec., p. 647.

Hensel, Harvey L. — Food Package Labeling—Legal Requirements and Commercial Practices. Aug., p. 392.

Highlights of the Report on State and Local Food and Drug Programs. Jan., p. 13.

Hoge, James F.
Factory Inspection Under the Federal Food, Drug and Cosmetic Act (Section 704). Dec., p. 673.
A Lawyer's View of Drug Investigation. Apr., p. 184.
Rendezvous with Destiny. Aug., p. 431.

I

International drug reporting system
Adoption by WHO. June, p. 322.

Introductory Statement (Twenty-first Annual Meeting of the Section on Food, Drug and Cosmetic Law of the New York State Bar Association). Feb., p. 109.

Investigational Drug Branch; Intra-FDA Relationships. Feb., p. 102.

K

Karl, Paul M.—The Advantages of, and Need for the Establishment of Uniform Guiding Principles and Model Standards for Food. Dec., p. 680.

Kefauver-Harris Amendments of 1962
Regulations issued by FDA. Jan., p. 28.
Three Years Later. Jan., p. 21.

Kelly, John T.—Three Years Later. Jan., p. 21.

Kelsey, Frances O. — Investigational Drug Branch: Intra-FDA Relationships. Feb., p. 102.

Kennedy, Cornelius B.—Now That the Fair Packaging and Labeling Act Is Law. Dec., p. 632.

L

Labeling
Food Package Labeling — Legal Requirements and Commercial Practices. Aug., p. 392.

Latin-American Food Code
Chapter VII. June, p. 312.
Chapter XVIII. Aug., p. 404.

Law Governing FDA Factory Inspection (The). May, p. 275.

Lawyer's View of Drug Investigation (A). Apr., p. 184.

Legislation
President's Message on Consumer Interest. Mar., p. 179.
Uniformity. Mar., p. 145.

Litchfield, John — Clinical Evaluation of Drug Safety. June, p. 348.

Look at FDA's New Rules of Practice —and Problems Still Unsolved (A). Sept., p. 448.

M

Manufacturing practice
Federal Food, Drug & Cosmetic Act, Sec. 501(a)(2)(B). Mar., p. 137.

McFarlane, Alexander N. — Opening Remarks (FLI — FDA's Ninth Annual Educational Conference). Jan., p. 4.

McMurray, R. D.—Some Recent Legal Developments in the Food, Drug and Cosmetic Field. Oct., p. 517.

McNally, Elaine G.—What the Consumer Expects, and Receives, from the Administration of the Federal Food, Drug and Cosmetic Act. July, p. 377.

Model Weights and Measures Law
Packaging Industry. Aug., p. 396.

More on "Zero Tolerance." May, p. 305.

Myers, Maven J. — Physician Owned Pharmaceutical Companies—A Wrong Without a Remedy. Aug., p. 412.

N

New drugs

Clinical evaluation. June, p. 348.
FDA's opinion that a new drug will become an old drug. June, p. 322.
Investigational procedures. Feb., p. 102.
Legal procedures. Feb., p. 68.

No Residue

Proposed Alternative to Zero Level and No Residue Regulations. Feb., p. 124.

Now That the Fair Packaging and Labeling Act is Law. Dec., p. 632.

O

O'Keefe, Harold F.

Compliance with Drug Abuse Control Amendments of 1965. July, p. 360.
Food and Drug Administration Industry Information Programs. Jan., p. 52.

Opening Remarks (FLI—FDA's Ninth Annual Educational Conference). Jan., p. 4.

Oser, Bernard L.

More on "Zero Tolerance." May, p. 305.
The Proposed Alternative to Zero Level and No Residue Regulations. Feb., p. 124.
The Scientists' Forum: Introduction to Food Additives. Nov., p. 616.

P

Packaging Industries and the Food Additives Amendment of 1958—It's Time for a Change in the Law (The). Dec., p. 647.

Packaging Industry Looks at the Model Law and Regulation (The). Aug., p. 396.

Pendergast, W. R.—Some Recent Legal Developments in the Food, Drug and Cosmetic Field. Oct., p. 517.

Peterson, Esther—The Consumer's Interest. Feb., p. 92.

Pharmaceutical Industry

Appraisal of progress. Jan., p. 33.
"Generic every time" case. Apr., p. 226.
Physician owned. Aug., p. 412.

Physician Owned Pharmaceutical Companies—A Wrong Without a Remedy. Aug., p. 412.

Pisani, Joseph M.—Drug Safety and the FDA. Feb., p. 68.

Prescription Drugs

Principles of warranty, failure to warn and strict liability in tort. Nov., p. 599.

Prescription or Ethical Drugs: Fallacies as to Warranties, Failure to Warn, and Strict Liability in Tort. Nov., p. 599.

President's Message on Consumer Interests (The). Mar., p. 179.

Product Liability

Federal government's position. June, p. 322.
1965. Mar., p. 154.
Prescription of Ethical Drugs: Fallacies as to Warranties, Failure to Warn and Strict Liability in Tort. Nov., p. 599.

Product Liability—1965. Mar., p. 154.

Progress in Consumer Education, Feb., p. 78.

Progress in Research — A Question. Jan., p. 28.

Proposed Alternative to Zero Level and No Residue Regulations (The). Feb., p. 124.

Proprietary Drug Industry

Federal regulation. Aug., p. 431.

Q

Question and Answer Panel of the FDA — FLI Ninth Annual Educational Conference. June, p. 322.

R

Rendezvous with Destiny. Aug., p. 431.

Report of the Third Session of the Joint FAO/WHO Codex Alimentarius Commission. Mar., p. 168.

Reports to the Reader. Jan., p. 3; Feb., p. 67; Mar., p. 135; Apr., p. 183; May, p. 247; June, p. 311; July, p. 359; Aug., p. 391; Sept., p. 439; Oct., p. 487; Nov., p. 551, Dec., p. 631.

Research

FDA research program. June, p. 322.

Food Research and the Food Laws. Apr., p. 191.

Progress in Research—A Question. Jan., p. 28.

Ricketts, E. F.—Highlights of the Report on State and Local Food and Drug Programs. Jan., p. 13.

S

Salmonella

FDA's position. June, p. 322.

Sanders, Howard J. The Scientists' Forum: Food Additives. Nov., p. 616.

Sayer, Murray D.—Artificial Sweeteners —Their Impact on the Food Laws. Feb., p. 111.

Scientists' Forum (The)
Food Additives. Nov., p. 616.

Some Recent Legal Developments in the Food, Drug and Cosmetic Field. Oct., p. 517.

Spiker, Earl G. — A Look at FDA's New Rules of Practice—and Problems Still Unsolved. Sept., p. 448.

Stafford, P. Gordon—A Look at FDA's New Rules of Practice—and Problems Still Unsolved. Sept., p. 448.

Standards of identity

Breaded shrimp concept. June, p. 322.

Sweeney, Harry A., Jr.—The "Generic Every Time" Case: Prescription Drug Industry *in Extremis*. Apr., p. 226.

T

Three Years Later. Jan., p. 21.

Toxicologic Aspects of Drug Safety. June, p. 336.

Trawick, James L.—Progress in Consumer Education. Feb., p. 78.

U

Uniformity—A Legislator's Viewpoint. Mar., p. 145.

V

Ventura, Sergio—The Common Market and Harmonization of the Food Laws. Sept., p. 440.

Vitamin D

Safety. June, p. 322.

Voluntary compliance

Food industry. Jan., p. 6.
Food research. Apr., p. 191.

W

Warren, William C. — Congressional Investigations; Some Observations. Jan., p. 40.

What the Consumer Expects, and Receives, from the Administration of the Federal Food, Drug and Cosmetic Act. July, p. 377.

What the Food Industry Needs from the FDA. July, p. 373.

Wolf, Ann M., Translator—Latin-American Food Code. June, p. 312; Aug., p. 404.

Z

Zero tolerance

FDA's position. June, p. 322.

More on "Zero Tolerance." May, p. 305.

Proposed Alternative to Zero Level and No Residue Regulations. Feb., p. 124.

Zero penicillin cross-examination. June, p. 322.